

This Page Is Inserted by IFW Operations
and is not a part of the Official Record

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

- BLACK BORDERS
- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

IMAGES ARE BEST AVAILABLE COPY.

**As rescanning documents *will not* correct images,
please do not report the images to the
Image Problem Mailbox.**



PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification⁴ : A61K 37/36	A1	(11) International Publication Number: WO 89/11293 (43) International Publication Date: 30 November 1989 (30.11.89)
(21) International Application Number: PCT/US89/02229 (22) International Filing Date: 22 May 1989 (22.05.89) (30) Priority data: 196,975 20 May 1988 (20.05.88) US (71) Applicants: INSTITUTE OF MOLECULAR BIOLOGY, INC. [US/US]; 812 Huntington Avenue, Boston, MA 02115 (US). PRESIDENT AND FELLOWS OF HARVARD COLLEGE [US/US]; 17 Quincy Street, Cambridge, MA 02138 (US). (72) Inventors: ANTONIAOES, Harry, N. ; 21 Magnolia Drive, Newton, MA 02148 (US). LYNCH, Samuel, E. ; 224 Jamaica Way, Jamaica Plain, MA 02158 (US). (74) Agent: CLARK, Paul, T.; Fish & Richardson, One Financial Center, Suite 2500, Boston, MA 02111-2658 (US).		(81) Designated States: AT (European patent), BE (European patent), CH (European patent), DE (European patent), FR (European patent), GB (European patent), IT (European patent), JP, LU (European patent), NL (European patent), SE (European patent). Published <i>With international search report.</i>
(54) Title: WOUND HEALING (57) Abstract Healing an external wound of a mammal by administering to the mammal a composition containing purified insulin-like growth factor-1 and purified transforming growth factor beta.		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	FI	Finland	ML	Mali
AU	Australia	FR	France	MR	Mauritania
BB	Barbados	GA	Gabon	MW	Malawi
BE	Belgium	GB	United Kingdom	NL	Netherlands
BF	Burkina Faso	HU	Hungary	NO	Norway
BG	Bulgaria	IT	Italy	RO	Romania
BJ	Benin	JP	Japan	SD	Sudan
BR	Brazil	KP	Democratic People's Republic of Korea	SE	Sweden
CF	Central African Republic	KR	Republic of Korea	SN	Senegal
CG	Congo	LI	Liechtenstein	SU	Soviet Union
CH	Switzerland	LK	Sri Lanka	TD	Chad
CM	Cameroon	LU	Luxembourg	TG	Togo
DE	Germany, Federal Republic of	MC	Monaco	US	United States of America
DK	Denmark	MG	Madagascar		
ES	Spain				

- 1 -
WOUND HEALING

Background of the Invention

This invention relates to healing wounds.

Growth factors are polypeptide hormones which
5 stimulate a defined population of target cells.
Examples of growth factors include platelet-derived
growth factor (PDGF), insulin-like growth factor
(IGF-I), transforming growth factor beta (TGF- β),
transforming growth factor alpha (TGF- α), epidermal
10 growth factor (EGF), and fibroblast growth factor
(FGF).

TGF- β is a multifunctional regulatory
polypeptide synthesized by many cell types and
sequestered in human platelets in amounts similar to
15 PDGF. The in vitro biological effects of TGF- β are
dependent upon the presence of other growth factors:
TGF- β in the presence of PDGF stimulates fibroblast
growth, and in the presence of EGF inhibits fibroblasts
(Roberts, et al. Proc. Natl. Acad. Sci. USA 82:119).
20 TGF- β inhibits proliferation of epithelial cells in
vitro (Shipley et al., 1986, Cancer Res. 46:2068), and
in vivo stimulates DNA, total protein, and collagen
synthesis when injected into wound chambers (Sporn et
al., 1986, Science 219:1329). The breaking strength of
25 incisional wounds increases in a dose dependent manner
after application of TGF- β (Mustoe, et al., Science
237:1333).

IGF-1 is synthesized de novo in the liver and
secreted into the plasma. In vitro, IGF-1 can promote
30 DNA synthesis in both mesenchymal and epithelial cells
(Van Wyk 1984, Hormonal Proteins and Peptides, Li,
ed.). Addition of IGF-1 in vivo by itself does not
promote wound healing, but when added with PDGF the
combination stimulates connective tissue and epithelial

- 2 -

proliferation (Lynch, et al., 1987, Proc. Natl. Acad. Sci., USA 84:7696).

Summary of the Invention

5 In general, the invention features healing an external wound in a mammal, e.g., a human patient, by applying to the wound an effective amount of a composition that includes a combination of purified TGF- β and purified IGF-1. Preferably, the TGF- β is human TGF- β , but can also be of another mammalian
10 species, e.g., porcine. The TGF- β can be isolated from natural sources (e.g., platelets) or, more preferably, produced by recombinant cells. The IGF-1 can be produced using recombinant DNA technology or solid phase peptide synthesis.

15 The composition of the invention aids in healing the wound, at least in part, by promoting the growth of epithelial and connective tissue and the synthesis of total protein and collagen. Wound healing using the composition of the invention is more effective
20 than that achieved in the absence of treatment (i.e., without applying exogenous agents) or by treatment with purified IGF-1 alone, or purified TGF- β alone.

In preferred embodiments of the invention, the composition is prepared by combining, in a
25 pharmaceutically acceptable carrier substance, e.g., commercially available inert gels or liquids (e.g., saline supplemented with albumin or methyl cellulose), purified IGF-1 and TGF- β (both of which are commercially available). Most preferably purified IGF-1 and TGF- β
30 are combined in a weight-to-weight ratio of between 1:4 and 4:1, preferably between 1:2 and 2:1, and more preferably 1:1 or 2:1. The purified TGF- β may be obtained from human platelets or by recombinant DNA technology. Thus, by the term "TGF- β " we mean both

- 3 -

platelet-derived and recombinant materials of mammalian, preferably primate, origin; most preferably, the primate is a human, but can also be a chimpanzee or other primate. Recombinant TGF- β can be recombinant monomer or homodimer, made by inserting into cultured prokaryotic or eukaryotic cells a DNA sequence encoding a subunit, and then allowing the translated subunits to be processed by the cells to form a homodimer.

The term "purified" as used herein refers to IGF-1 or TGF- β which, prior to mixing with the other, is 95% or greater, by weight, IGF-1 or TGF- β , i.e., is substantially free of other proteins, lipids, and carbohydrates with which it is naturally associated.

A purified protein preparation will generally yield a single major band on a polyacrylamide gel for each IGF-1 or TGF- β component. Most preferably, the purified IGF-1 or TGF- β used in the composition of the invention is pure as judged by amino-terminal amino acid sequence analysis.

The composition of the invention provides a fast, effective method for healing external wounds of mammals, e.g., bed sores, lacerations and burns. The composition enhances connective tissue formation compared to natural healing (i.e., with no exogenous agents added) or pure IGF-1 or TGF- β alone. Unlike pure TGF- β alone, the composition promotes a significant increase in both new epithelial tissue and connective tissue. The epithelial layer obtained is thicker than that created by natural healing or by TGF- β alone, and also contains more epithelial projections connecting it to the new connective tissue; it is thus more firmly bound and protective.

Other features and advantages of the invention will be apparent from the following description of the

- 4 -

preferred embodiments thereof, and from the claims.

Description of the Preferred Embodiments

We now describe preferred embodiments of the invention.

5 External wounds, e.g., bed sores and burns, are treated, according to the invention, with IGF-1/TGF- β . Recombinant human IGF-1 is commercially available from Amgen Biologicals (Thousand Oaks, CA). Purified human and porcine TGF- β are commercially available from R&D
10 Systems, Inc. (Minnesota, MN).

 TGF- β was purified from human or porcine platelets by the method of Assoian (1983, J. Biol. Chem. 258: 7155). Briefly, platelet-rich plasma (20-30 units, 2-5 days old) was centrifuged (3200xg 30 min, at 4°C) to
15 remove plasma proteins. The platelets were then washed twice in 500ml portions of Tris-HCl/citrate buffer, and recentrifuged. The washed platelets were then added to a solution of acid ethanol and then immediately extracted in a homogenizer. After incubation overnight
20 at 4°C, precipitated proteins were removed by centrifugation and the supernatant adjusted to pH 3 using NH_4OH . TGF- β was precipitated by addition of ethanol (2 volumes at 0°C) and ethyl ether (4 volumes at 0°C). The precipitate was collected by centrifugation
25 and suspended in 1M acetic acid (10ml). The supernatant was separated from the precipitate by centrifugation and placed on a Bio-Gel P60 gel filtration column (4.4 x 115 cm), with a flow rate of 20ml/hr, equilibrated in 1M acetic acid. Five milliliter fractions were collected
30 and assayed for biological activity using growth inhibition of BALB/MK cells and anchorage-independent growth of non-neoplastic NRK fibroblasts. Fractions containing peak activity were pooled, lyophilized, and redissolved in 0.5ml of 1M acetic acid containing 8M

- 5 -

ultra-pure urea (Schwartz/Mann) and gel filtered at a flow rate of 3ml/hr on a Bio-Gel P60 column (1.6 x 85cm). Aliquots of column fractions were tested for TGF- β activity as described above. Fractions containing peak TGF- β activity were pooled, dialyzed against 1M acetic acid to remove urea, and added to a C-18 (synchropak) HPLC column in 0.1% trifluoroacetic acid and eluted with a 20-50% acetonitrile gradient. Biologically active fractions were pooled, and final purity checked by SDS-PAGE and amino acid analysis for known properties of TGF- β .

Recombinant TGF- β can be prepared by standard techniques. For example, oligonucleotide probes designed on the basis of the protein sequence of TGF- β can be used for the isolation of TGF- β exons in a human genomic DNA or a cDNA library, using the technique described in Birnbaech (1985, Nature 316: 701). The gene for TGF- β is isolated, cloned into standard expression vectors, and transfected into mammalian cells, from which TGF- β is then purified using standard methods.

Wound Healing

To determine the effectiveness of IGF-1/TGF- β mixtures in promoting wound healing, the following experiments were performed.

Young white Yorkshire pigs (Parson's Farm, Hadley, MA) weighing between 10 and 15 kg were fasted for at least 6 hours prior to surgery and then anesthetized. Under aseptic conditions, the back and thoracic areas were clipped, shaved, and washed with mild soap and water. The area to be wounded was then disinfected with 70% alcohol.

Wounds measuring 1 cm x 2 cm were induced at a depth of 0.5 mm using a modified Castroviejo electrokeratome (Storz, St. Louis, MO, as modified by

- 6 -

Brownells, Inc.). The wounds resulted in complete removal of the epithelium, as well as a portion of the underlying dermis (comparable to a second degree burn injury). Individual wounds were separated by at least 15 mm of unwounded skin. Wounds receiving identical treatment were organized as a group and separated from other groups by at least 3 cm. Wounds receiving no growth factor treatment were separated from wounds receiving such treatment by at least 10 cm.

The wounds were treated directly with a single application of the following growth factors suspended in biocompatible gel: 1) 500 ng pure human or porcine TGF- β ; 2) 500 ng pure recombinant IGF-1 alone; 3) 500 ng human or porcine TGF- β plus 500 ng pure recombinant IGF-1.

Following wounding, biopsy specimens were taken on days 3 through 10. Biopsy specimens for histologic evaluation were taken as wedges approximately 3 mm deep and placed in 10% formalin. Specimens for biochemical analysis were obtained using an electrokeratome. The final dimensions of the specimens were 1.5 mm x 10 mm x 1.5 mm. Three specimens per wound were collected for biochemical analysis. Following collection, the specimens were frozen in liquid Nitrogen and stored at -80°C.

Histologic Evaluation

Histologic specimens were prepared using standard paraffin impregnating and embedding techniques. Four micron sections were made and stained using filtered Harris hematoxylin and alcoholic eosin; they were then observed under a microscope. Computer-aided morphometric analyses were performed. The area of the new epithelial and connective tissue layers were assessed with the aid of a customized

- 7 -

program (need details) for determining areas of histological specimens.

Collagen Determination

The specimens for biochemical analysis were thawed and the newly synthesized wound tissue dissected from the surrounding tissue under a dissecting microscope. The samples were hydrolyzed in 6M HCl at 120°C for 18 hours in sealed ampoules. Assay of the hydrolysate for hydroxyproline, an amino acid unique to collagen was then performed using the technique of Switzer and Summer, 1971, Anal. Biochem. 39:487.

Results

The results from histologic evaluation indicated that wounds treated with TGF-β had a thinner epithelial layer than wounds receiving no treatment. In contrast, wounds treated with the combination of purified human or porcine TGF-β and recombinant human IGF-1 had thicker connective tissue and epithelial layers, and more extensive epithelial projections connecting these layers, than wounds receiving no treatment, human or porcine TGF-β alone, or pure IGF-1 alone. The IGF-1 plus TGF-β-treated wounds also had greater total collagen content, as indicated by increased hydroxyproline, than wounds treated with TGF-β alone, IGF-1 alone, or gel alone.

Dosage

To determine the appropriate dosage of purified TGF-β, the above-described experiments were repeated except that the wounds were treated with 2.5 ng, 5.0 ng, and 10 ng of purified TGF-β per square millimeter of wound dispersed in 30μl of biocompatible gel. The results showed that optimum effects were produced when the TGF-β content of a IGF-1/TGF-β mixture was 5.0 ng/mm² or higher.

- 8 -

To determine the optimal ratio of IGF-1 to TGF- β , combinations in which the weight to weight ratio of IGF-1 to TGF- β ranged from 1:10 to 25:1 were evaluated as described above. Optimum results were achieved with a ratio of between 1:2 and 2:1.

Other Embodiments

Other embodiments are within the following claims. For example, IGF-1 and TGF- β can be obtained by standard recombinant DNA technology using nucleic acid having a base sequence identical to that of the naturally occurring gene encoding IGF-1 or TGF- β in a human or other mammal. Further, this nucleic acid may be modified by conservative base substitutions such that it encodes the same amino acid sequence of naturally occurring IGF-1 or TGF- β ; or modified with base substitutions which encode a different amino acid sequence to that naturally occurring, but the protein product of which has substantially the same wound healing properties as the naturally occurring proteins.

- 9 -
Claims

1. A method for healing an external wound of a mammal comprising applying to said wound a wound-healing amount of a composition comprising purified Insulin-like growth factor-1 and purified transforming growth factor beta.
2. The method of claim 1 wherein the weight to weight ratio of said Insulin-like growth factor-1 to said transforming growth factor beta in said composition is between 1:4 and 25:1.
3. The method of claim 2 wherein said ratio is between 1:2 and 10:1.
4. The method of claim 3 wherein said ratio is about 1:2 or 2:1.
5. A wound healing composition comprising purified Insulin-like growth factor-1 and purified transforming growth factor beta, in a weight to weight ratio of 1:4 to 25:1.
6. The composition of claim 5 wherein said ratio is between 1:2 and 10:1.
7. The composition of claim 6 wherein said ratio is about 1:2 or 2:1.
8. A method for preparing a composition for healing wounds, comprising mixing purified Insulin-like growth factor-1 and purified transforming growth factor beta in a weight to weight ratio of between 1:4 and 25:1.

INTERNATIONAL SEARCH REPORT

International Application No PCT/US89/02229

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) ⁶ According to International Patent Classification (IPC) or to both National Classification and IPC IPC (4): A61K 37/36 U.S. CL: 514/8,12,21		
II. FIELDS SEARCHED Minimum Documentation Searched ⁷		
Classification System	Classification Symbols	
US	514/8,12,21	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched ⁸		
III. DOCUMENTS CONSIDERED TO BE RELEVANT ⁹		
Category ¹⁰	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³
X	W, U.S. Application Serial Number 468,590, Sporn, Published 1983 See pages 24-25	1-8
<p> ¹⁰ Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. "Δ" document member of the same patent family </p>		
IV. CERTIFICATION		
Date of the Actual Completion of the International Search 14 August 1989 International Searching Authority ISA/US		Date of Mailing of this International Search Report 15 SEP 1989 Signature of Authorized Officer Howard E. Schain